

INFORMATION DISCLOSURE
CITATION

ATTY. DOCKET NO.

1430-252

SERIAL NO.

09/646,224

APPLICANT

GROSE et al.

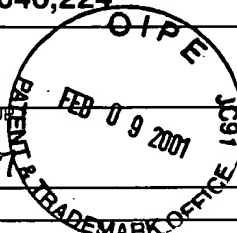
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September 14, 2000

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1647



U.S. PATENT DOCUMENTS

*EXAMINER INITIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE

OTHER DOCUMENTS (including Author, Title, Date, Pertinent pages, etc.)

BE	14	V58419; WO 98/38302-A2; 03 Sep 1998.
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Sheet 2 of 2



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<input checked="" type="checkbox"/>	Ptacek et al, "Identification of a mutation in the gene causing hyperkalemic periodic paralysis", Cell 1991 Nov 29;67(5):1021-7.
<input checked="" type="checkbox"/>	Ptacek et al, "Mutations in an S4 segment of the adult skeletal muscle sodium channel cause paramyotonia congenita", Neuron 1992 May;8(5):891-7.
<input checked="" type="checkbox"/>	McClatchey et al, "Novel mutations in families with unusual and variable disorders of the skeletal muscle sodium channel", Nat Genet 1992 Oct;2(2):148-52.
<input checked="" type="checkbox"/>	Lerche et al, "Human sodium channel myotonia: slowed channel inactivation due to substitutions for a glycine within the III-IV linker", J Physiol (Lond) 1993 Oct;470:13-22.
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<input checked="" type="checkbox"/>	Heine et al, "A novel SCN4A mutation causing myotonia aggravated by cold and potassium", Hum Mol Genet 1993 Sep;2(9):1349-53.
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